California Drug Recall Information

Recall Name
PharMEDium Services Recalls Norepinephrine Bitartrate Added to 0.9% Sodium Chloride Due to Discoloration in the Admixture

Recall Date	Product Description	Recalling Firm	Recall Reason
12/22/15	• 4mg Norepinephrine Bitartrate (16mcg/mL) added to 0.9% Sodium Chloride in 250mL Viaflex Bag **NDC # 61553-134-61** • 8mg Norepinephrine Bitartrate (32mcg/mL) added to 0.9% Sodium Chloride in 250mL Viaflex Bag **NDC # 61553-127-61**	**PharMEDium Services, LLC** Lake Forest, IL	*Due to complaints of discoloration.* *Discoloration is indicative of degradation and could result in decreased potency due to oxidation of Norepinephrine Bitartrate.*

Recall Class	Product Identification	Distribution	Affected Dates
N/A	Click this link for affected lots.	**CA**, nationwide	Expiration Date: 02/10/16 to 02/18/16

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm479677.htm